## WHAT IS CLAIMED IS:

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A method of refolding a first insoluble, recombinant, eukaryotic 1. 1 glycosyltransferase, wherein the glycosyltransferase comprises a maltose binding protein 2 domain (MBD), the method comprising the steps of 3 (a) solubilizing the insoluble, recombinant, eukaryotic glycosyltransferase in a 4 5 solubilization buffer; and (b) contacting the soluble eukaryotic glycosyltransferase with a refolding 6 buffer comprising a redox couple to refold the eukaryotic glycosyltransferase, wherein the 7 refolded eukaryotic glycosyltransferase catalyzes the transfer of a sugar from a donor 8 substrate to an acceptor substrate. 9

- 1 2. The method of claim 1, wherein the first eukaryotic glycosyltransferase 2 is truncated to remove all or a portion of a stem region.
  - 3. The method of claim 1, wherein an unpaired cysteine in the first eukaryotic glycosyltransferase is removed by substitution with a non-cysteine amino acid.
  - 4. The method of claim 1, wherein the first eukaryotic glycosyltransferase is selected from the group consisting of GnT1, GalT1, StIII Gal3, St3GalI, St6 GalNAcTI, Core GalITI, GalNAcT2.
  - 5. The method of claim 1, wherein the first eukaryotic glycosyltransferase further comprises a purification domain selected from the group consisting of a starch binding domain, a thioredoxin domain, a SUMO domain, a poly-His domain, a myc epitope domain, and a glutathione-S-transferase domain.
- 1 6. The method of claim 1, wherein the first eukaryotic glycosyltransferase 2 further comprises a self cleaving domain.
- 7. The method of claim 1, wherein the first eukaryotic glycosyltransferase is expressed in a bacterial host cell as an insoluble inclusion body.
- 1 8. The method of claim 1, wherein a second insoluble, recombinant 2 eukaryotic glycosyltransferase is refolded with the first eukaryotic glycosyltransferase.

1	9. The method of claim 8, wherein a third insoluble, recombinant
2	eukaryotic glycosyltransferase is refolded with the first eukaryotic glycosyltransferase and
3	the second eukaryotic glycosyltransferase.
1	10. The method of claim 1, wherein the redox couple is selected from the
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2	group consisting of reduced glutathione/oxidized glutathione (GSH/GSSG) and cysteine/
3	cystamine.
1	11. The method of claim 1, wherein the acceptor substrate is selected from
2	a protein, a peptide, a glycoprotein, and a glycopeptide.
1	12. The method of claim 1, wherein the first eukaryotic glycosyltransferase
2	is a sialyltransferase.
1	13. The method of claim 12, wherein the sialyltransferase is selected from
2	the group consisting of StIII Gal3, St3GalI, St6 GalNAcTI.
1	14. The method of claim 12, wherein the donor substrate is a CMP-sialic
2	acid PEG molecule and the acceptor substrate is selected from a protein, a peptide, a
3	glycoprotein, and a glycopeptide.
1	15. A recombinant, eukaryotic glycosyltransferase, wherein a stem anchor
2	region and a transmembrane domain are deleted from the recombinant, eukaryotic
3	glycosyltransferase, and wherein the glycosyltransferase is fused in frame to a maltose
4	binding domain.
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1	16. The recombinant, eukaryotic glycosyltransferase of claim 15, wherein
2	all or a portion of the stem region is deleted.
1	17. The recombinant, eukaryotic glycosyltransferase of claim 15, wherein
2	an unpaired cysteine in the recombinant, eukaryotic glycosyltransferase is removed by
3	substitution with a non-cysteine amino acid.
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1	18. The recombinant, eukaryotic glycosyltransferase of claim 15, wherein
2	the glycosyltransferase is selected from the group consisting of a GnT1 protein, a GalT1
3	protein, an StIII Gal3 protein, an St3GalI protein, an St6 GalNAcTI protein, a Core GalITI
4	protein, and a GalNAcT2 protein.

1 19. The recombinant, eukaryotic glycosyltransferase of claim 15, wherein 2 the glycosyltransferase is a GnT1 protein.

- 1 20. The GnT1 protein of claim 19, wherein the GnT1 protein is a truncated 2 human GnT1 protein selected from GnT1 Δ35 and GnT1Δ103.
- 1 21. The GnT1 protein of claim 19, wherein the GnT1 protein is a human 2 GnT1 protein comprising an unpaired cysteine substitution selected from the group consisting
- 3 of CYS121ALA, CYS121ASP, and ARG120ALA, CYS121HIS.
- 1 22. The recombinant, eukaryotic glycosyltransferase of claim 15, wherein 2 the glycosyltransferase is a GalT1 protein.
- 1 23. The GalT1 protein of claim 22, wherein the GalT1 protein is a
- 2 truncated bovine GalT1 protein selected from GalT1  $\Delta$ 70 and GalT1  $\Delta$ 129.
- 1 24. The GalT1 protein of claim 22, wherein the GalT1 protein is a bovine 2 GalT1 protein comprising an unpaired cysteine substitution of CYS342THR.
- 1 25. The recombinant, eukaryotic glycosyltransferase of claim 15, wherein 2 the glycosyltransferase is an ST3GalIII protein.
- 26. The ST3GalIII protein of claim 25, wherein the ST3GalIII protein is a
  truncated rat ST3GalIII protein selected from ST3GalIII Δ28, ST3GalIII Δ73, ST3GalIII Δ85
  and ST3GalIII Δ86.
- 1 27. The recombinant, eukaryotic glycosyltransferase of claim 15, wherein 2 the glycosyltransferase is a Corel GalT1 protein.
- 1 28. The recombinant, eukaryotic glycosyltransferase of claim 15, wherein 2 the glycosyltransferase is an ST3Gal1 protein.
- 29. The ST3Gal1 protein of claim 28, wherein the ST3Gal1 protein is a
  truncated human ST3Gal1protein selected from ST3Gal1 Δ29, ST3Gal1 Δ45, and ST3Gal1
  Δ56.
- 1 30. The recombinant, eukaryotic glycosyltransferase of claim 15, wherein 2 the glycosyltransferase is an ST6GalNAc1 protein.

1 31. The recombinant, eukaryotic glycosyltransferase of claim 15, wherein 2 the glycosyltransferase is an GalNAcT2 protein.

- 1 32. The GalNAcT2 protein of claim 31, wherein the GalNAcT2 protein is
- 2 a truncated human GalNAcT2 protein selected from GalNAcT2  $\Delta$ 40, GalNAcT2  $\Delta$ 51,
- 3 GalNAcT2  $\Delta$ 74 and GalNAcT2  $\Delta$ 95.
- 1 33. A method of remodeling a protein, a peptide, a glycoprotein, or a
- 2 glycopeptide using the recombinant, eukaryotic glycosyltransferase of claim 15.

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